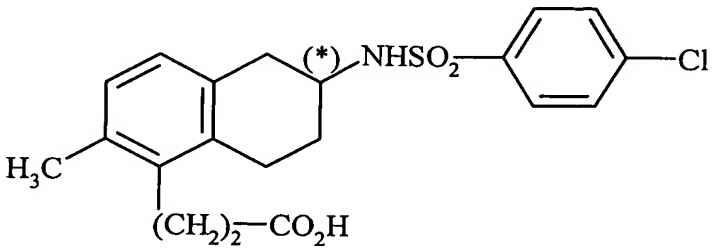
CLAIMS

1- Solid orodispersible pharmaceutical composition of compound A of formula (I), optionally in the form of an optical isomer, or a pharmaceutically acceptable salt thereof :



(I),

characterised in that it comprises :

- compound A or a pharmaceutically acceptable salt thereof,
- granules consisting of co-dried lactose and starch.

2- Pharmaceutical composition according to claim 1, characterised in that compound A is in the form of an optical isomer of configuration (R).

3- Pharmaceutical composition according to either claim 1 or claim 2, characterised in that it comprises, in relation to the total weight of the composition :

- from 2.5 % to 20 % by weight of compound A or a pharmaceutically acceptable salt thereof,
- from 75 % to 95 % by weight of granules consisting of co-dried lactose and starch.

4- Pharmaceutical composition according to claim 3, characterised in that it comprises from 5 % to 10 % by weight of compound A or a pharmaceutically acceptable salt thereof.

5- Pharmaceutical composition according to any one of claims 1 to 4, characterised in that compound A is in the form of a sodium salt.

6- Pharmaceutical composition according to claim 1, characterised in that it also comprises one or more flavourings, and sweeteners.

7- Pharmaceutical composition according to claim 1, characterised in that it also comprises one or more lubricants and a flow agent.

8- Pharmaceutical composition according to any one of claims 1 to 7, characterised in that it is in the form of a tablet.

9- Tablet according to claim 8, characterised in that it is obtained by direct compression.

10- Tablet according to claim 9, characterised in that its hardness is from 15 to 30 Newtons.

11- Use of granules consisting of co-dried lactose and starch in the manufacture of solid orodispersible compositions of compound A, which disintegrate in the mouth in less than three minutes, preferably less than one minute, for oral or buccal administration.

12- Solid orodispersible pharmaceutical composition of compound A according to claim 1, or a pharmaceutically acceptable salt thereof, for obtaining an antithrombotic medicament.